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Pfizer conducted a report entitled: CUMULATIVE ANALYSIS OF POST-AUTHORIZATION ADVERSE EVENT REPORTS OF PF-07302048 (BNT162B2) RECEIVED THROUGH 28-FEB-2021 The objective of the Classified Pfizer Report was to tabulate and categorize detailed data pertaining to the impacts of the Pfizer BioNTech mRNA vaccine on, Mortality, Morbidity and Adverse Events from December 14, 2020 to February 28, 2021.

The declassified data provides a stark overview of the vaccine's impact during the first 90 days of its global release.The company received a cumulative total of 42,086 case reports. Within these reports, there were 158,893 individual adverse events documented.

Some significant Categories

- Respiratory, Thoracic, and Mediastinal Disorders: 8,848 events.
- Skin and Subcutaneous Tissue Disorders: 8,476 events.
- Injury, Poisoning, and Procedural Complications: 5,590 events.
- Infections and Infestations: 4,610 events.
- Investigations: 3,693 events.